IN THE CLAIMS

This listing of claims replaces all prior versions, and listings, in this application.

1. (currently amended) A drug/gene eluting stent comprising a <u>surface</u> layer containing a gene encoding a hybrid polypeptide, <u>wherein the hybrid polypeptide comprises a fibronectin-derived collagen binding domain (FNCBD) polypeptide and an N-terminal deleted monocyte chemoattractant protein-1 (MCP-1) on the surface.</u>

Claims 2-4 (canceled)

- 5. (currently amended) The drug/gene eluting stent of claim 1 comprising:
- (i) a primer layer applied to an exterior surface of the stent,
- (ii) a drug layer base coated on the primer layer,
- (iii) a gene-containing layer formed by absorbing the gene in the drug layer base, and
- (iv) a protective layer coated on the gene-containing layer. The drug/gene eluting stent according to claim 4, wherein the N-terminal deleted chemokine is N-terminal deleted compound (7ND) of a monocyte chemoattractant protein-1 (MCP-1).
- 6. (currently amended) The drug/gene eluting stent of according to claim 1, wherein the gene encoding the hybrid polypeptide comprises has the nucleotide sequence shown showin in SEQ ID No: 1-or 2.

Claims 7-8 (canceled)

9. (currently amended) A method for treating vascular restenosis, acute coronary syndromes or cerebral ischemia, which comprises <u>placing a using the-drug/gene</u> eluting stent <u>comprising a surface layer which contains a gene encoding a hybrid polypeptide</u>, wherein the hybrid polypeptide comprises a fibronectin-derived collagen binding domain (FNCBD) polypeptide and an N-terminal deleted monocyte chemoattractant protein-1 (MCP-1) in a blood vessel-according to claim 1.

- 10. (currently amended) The method according to claim 9, wherein vascular restenosis is reduced as compared to placing a stent which does not contain a gene encoding a hybrid polypeptide comprising FNCBD and an N-terminal deleted MCP-1 in a blood vessel. Use of the drug/gene eluting stent according to claim 1 for manufacturing an agent for treating vascular restenosis, acute coronary syndromes or cerebral ischemia.
- 11. (new) The method according to claim 9, wherein the drug/gene eluting stent comprises:
- (i) a primer layer applied to an exterior surface of the stent,
- (ii) a drug layer base coated on the primer layer,
- (iii) a gene-containing layer formed by absorbing the gene in the drug layer base, and
- (iv) a protective layer coated on the gene-containing layer.
- 12. (new) The method according to claim 9, wherein the gene encoding the hybrid polypeptide comprises the nucleotide sequence shown in SEQ ID No: 1.
- 13. (new) The method according to claim 9, wherein vascular restenosis is treated.
- 14. (new) The method according to claim 13, wherein the vascular restenosis is a relapsed stenosis of post percutaneous transluminal coronary angioplasty (PTCA) or percutaneous transluminal angioplasty (PTA).
- 15. (new) The method according to claim 9 further comprising post percutaneous transluminal coronary angioplasty (PTCA) or percutaneous transluminal angioplasty (PTA) prior to placing the drug/gene eluting stent in the blood vessel.
- 16. (new and withdrawn) A drug/gene eluting stent comprising a surface layer containing a gene encoding a hybrid polypeptide, wherein the hybrid polypeptide

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comprises a fibronectin-derived collagen binding domain (FNCBD) polypeptide and an HGF.

- 17. (new and withdrawn) The drug/gene eluting stent of claim 16, wherein the gene encoding the hybrid polypeptide comprises the nucleotide sequence shown in SEQ ID No: 2.
- 18. (new and withdrawn) A method for treating vascular restenosis, acute coronary syndromes or cerebral ischemia, which comprises placing the drug/gene eluting stent of claims 17 in a blood vessel.
- 19. (new and withdrawn) The method according to claim 18, wherein the gene encoding the hybrid polypeptide comprises the nucleotide sequence shown in SEQ ID No: 2.